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UTILITY PATENT APPLICATION TRANSMITTAL

 Attorney
Docket No.

 96074DIV
(3600-011-01)

Total Pages

80

 Pay for new non-provisional applications under 37 CFR 1.53(b))
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First Named Inventor or Application Identifier

 Ranjan GHOSAL, James A. BELMONT, Douglas M. SMITH, and
 Janeel MENASHI

Express Mail Label No. EL352564703US

 APPLICATION ELEMENTS FOR:
 MODIFIED CARBON ADSORBENTS AND PROCESSES FOR
 ADSORPTION USING THE SAME

 ADDRESS TO:
 Assistant Commissioner for Patents
 BOX PATENT APPLICATIONS
 Washington, D.C. 20231

 575 U.S. PTO
 09/475385

12/30/99

 1. ☒ Fee Transmittal Form (Incorporated within this form) (Submit an original and a duplicate for fee processing)

 2. ☒ Application

Total Pages [32]

 3. ☒ Drawing(s) (35 USC 113) (Figs. 1-5)

Total Sheets [5]

 4. ☒ Oath or Declaration and Power of Attorney

Total Pages [12]

 a. ☐ Newly executed (original or copy)

 b. ☒ Copy from prior application (37 CFR 1.63(d) (for continuation/divisional with Box 17 completed).

 i. ☐ Deletion of Inventor(s) Signed statement attached deleting inventor(s) named in prior application, see 37 CFR 1.63(d)(2) and 1.33(b).

 5. ☒ Incorporation by reference (useable if box 4b is checked)

 The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied under
 Box 4b, is considered as being part of the disclosure of the accompanying application and is hereby incorporated by
 reference therein

 6. ☐ Microfiche Computer Program (Appendix)

 7. ☐ Nucleotide and/or Amino Acid Sequence Submission (if applicable, all necessary)

 a. ☐ Computer Readable Copy

 b. ☐ Paper Copy (identical to computer copy)

 c. ☐ Statement Verifying identity of above copies

ACCOMPANYING APPLICATION PARTS

 8. ☒ Assignment Papers (cover sheet and document(s))

Total Sheets [9]

 9. ☐ 37 CFR 3.73(b) Statement (when there is an assignee)

 Date December 30, 1999 Label No. EL352564703US

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UTILITY PATENT APPLICATION TRANSMITTAL

(Only for new non-provisional applications
under 37 CFR 1.53(b))

Attorney
Docket No.

970409DIV (3600-011-01)

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Janeel MENASHI

PAGE 2 OF 3

10. ☐ English translation Document (if applicable)

11. ☒ Information Disclosure Statement ☐ Copies of IDS Citations Total Pages [16]

12. ☒ Preliminary Amendment (3 PAGES) ☒ Please cancel claims 1-17, 26, and 27

13. ☒ Return Receipt Postcard (MPEP 503)

14. ☐ Small Entity Statement(s) ☐ Statement filed in prior application

15. ☐ Claim for Convention Priority ☐ Certified copy of Priority Document(s)

a. Priority of application no. filed on is claimed under 35 USC 119.

The certified copies/copy have/has been filed in prior application Serial No.

(For Continuing Applications, if applicable).

16. ☐ Other _____

17. If a CONTINUING APPLICATION, check appropriate box and supply the requisite information:

☐ Continuation ☐ Division ☐ Continuation-in-part (CIP).

FEE TRANSMITTAL	Number Filed	Number Extra	Rate	Basic Fee
The filing fee is calculated below				\$760.00
Total Claims	18 - 20	0	X \$18.00	\$000.00
Independent Claims	2 - 3	0	X \$78.00	\$000.00
Multiple Dependent Claims			\$260.00	
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Fee for recording enclosed Assignment			\$40.00	
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UTILITY PATENT
APPLICATION TRANSMITTAL

(Only for new non-provisional applications Under 37 CFR 1.53(b))

Attorney
Docket No.

970409DIV (3600-011-01)

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Janeel MENASHI

PAGE 3 OF 3

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18. CORRESPONDENCE ADDRESS

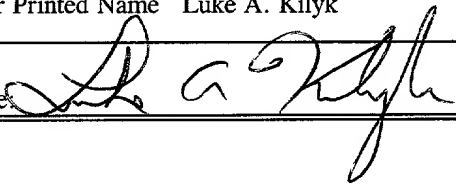
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Henry B. Smith

Name (Print)

Henry B. Smith

Signature

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of:)

Ranjan GHOSAL *et al.*)

Serial No.: Div. of 08/663,709)

Group Art Unit: Unknown

Filed: November 26, 1999)

Examiner: Unknown

For: MODIFIED CARBON ADSORBENTS AND PROCESSES
FOR ADSORPTION USING THE SAME

PRELIMINARY AMENDMENT

Assistant Commissioner for Patents
Washington, D.C. 20231

December 30, 1999

Sir:

Prior to examination of the above-identified new application, please amend the application as follows.

IN THE CLAIMS

Please cancel claims 1-17, 26, and 27. Please amend claim 18 as follows:

18. (Amended) A method to adsorb an adsorbate comprising contacting said adsorbate with a modified carbonaceous material capable of adsorbing said adsorbate wherein at least one organic group is attached to said modified carbonaceous material, wherein said organic group comprises an aromatic group or a C₁-C₁₂ alkyl group directly attached to the carbonaceous material.

Please add the following new claims.

--28. A catalyst comprising a modified carbonaceous material having attached at least one organic group, wherein said organic group comprises an aromatic group or a C₁-C₁₂ alkyl group directly attached to the carbonaceous material.

29. The catalyst of claim 28, wherein said organic group is (C₆H₄)-SO₃⁻Na⁺, (C₆H₄)-SO₃⁻Li⁺, OR (C₆H₄)SO₃⁻K⁺.

30. The catalyst of claim 28, wherein said organic group is p-C₆H₄SO₂NH₂ or -C₆H₄NH₂.

31. The catalyst of claim 28, wherein said organic group is hydrophilic.

32. The catalyst of claim 28, wherein said modified carbonaceous material is activated carbon or carbon black.

33. The catalyst of claim 28, wherein said organic group contains an exchangeable ion.

34. The catalyst of claim 28, wherein said organic group further comprises a functional group.

35. The catalyst of claim 28, further comprising an ionic group, an ionizable group, or a mixture thereof.

36. The catalyst of claim 34, wherein said functional group is a sulfonic acid group, a sulfinic acid group, a phosphonic acid group, a carboxylic acid group, a quaternary ammonium group, a quaternary phosphonium group, or a salt thereof.

Preliminary Amendment
U.S. Patent Application No. 08/663,709

37. The catalyst of claim 28, wherein said aromatic group is a phenyl group or a naphthyl group.--

REMARKS

Entry of this preliminary amendment is respectfully requested. Support for the new claims can be found throughout the specification, including the examples. Prompt and favorable consideration of the application is respectfully requested.

If there are any fees due in connection with the filing of this response, please charge the fees to deposit Account No. 03-0060. If a fee is required for an extension of time under 37 C.F.R. § 1.136 not accounted for above, such extension is requested and should also be charged to our Deposit Account.

Should the Examiner deem that any further action by applicants or applicants' undersigned representatives is desirable or required, the Examiner is invited to telephone the undersigned at the number set forth below.

Respectfully submitted,



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UNITED STATES PATENT APPLICATION
OF

RANJAN GHOSAL, JAMES A. BELMONT,
DOUGLAS M. SMITH AND JAMEEL MENASHI

FOR

MODIFIED CARBON ADSORBENTS AND PROCESSES FOR
ADSORPTION USING THE SAME

6502207 = 24 23 22 21 20 19 18 17 16 15 14 13 12 11 10 9 8 7 6 5 4 3 2 1

FIELD OF THE INVENTION

This invention relates to carbon adsorbents made from modified carbonaceous materials and also relates to a method of using these adsorbents, including a method to increase the adsorption capacity and/or alter the adsorption affinity of carbonaceous materials capable of adsorbing an adsorbate.

BACKGROUND OF THE INVENTION

Adsorption is an important operation in many industrial processes. The effectiveness of an adsorbent depends, primarily, on its surface area, pore structure, and surface chemistry. The nature of the adsorbate which is to be adsorbed frequently dictates the chemical nature of the adsorbent. For example, carbonaceous adsorbents are often used to selectively remove organic compounds from liquid, gaseous, or vapor media. Silica and alumina based adsorbents are employed to selectively adsorb polar adsorbates such as water, ammonia, and the like from similar media.

The efficacy of an adsorbent for a particular application is usually determined by the adsorption capacity and selectivity of the adsorbent for the adsorbate in question. The adsorption capacity may be measured per unit mass or per unit volume of the adsorbent. In general, the higher the adsorption capacity and selectivity of an adsorbent for a particular adsorbate, the more useful it is, since less of the adsorbent has to be used to effect the same removal of the adsorbate.

Carbonaceous materials, such as activated carbon, carbon black, and the like, represent an important class of adsorbents which are used in many fields such as separation, purification, and waste treatment, among others. Because of their widespread

use, any method for improving the adsorption properties of carbonaceous adsorbents for a particular adsorbate can have a large impact on the efficacy and economy of the processes utilizing them. Therefore, attempts have been made in the past to modify the surface chemistry of carbonaceous adsorbents. The methods employed for their modification can be broadly classified into physical and chemical means. In surface modification by physical means, a species is deposited on the surface of the carbonaceous adsorbent to form a layer which then changes its adsorption properties. However, such modification techniques have limited utility because the deposited layer is easily removed. In surface modification by chemical means, the modifying species is attached to the carbon surface by a chemical bonding mechanism.

The characteristics of the adsorption isotherm, representing the relationship between the extent of adsorption and adsorbate concentration or adsorbate partial pressure at a fixed temperature, is also of importance. As described by Sircar et al. in "Activated Carbon for Gas Separation and Storage," Carbon, Vol. 34, No. 1, pp. 1-12 (1996), the characteristics of the preferred adsorption isotherm will depend on the separation process being employed. For example, in cases where adsorbent regeneration is effected by a pressure swing, the preferred adsorbent is one with a moderate affinity for the adsorbate. When the adsorbate is strongly adsorbed, that is, when it has a strong affinity for the adsorbent, regeneration becomes difficult and energy intensive. On the other hand, when the adsorbent exhibits a weak affinity for the adsorbate, it has a small adsorption capacity at low adsorbent partial pressures and, hence, the adsorption mass transfer zone becomes

very long. Thus, the availability of a method for altering the affinity of an adsorbent for an adsorbate is advantageous.

Thus, any method for increasing the adsorption capacity and/or modifying the adsorption affinity of the adsorbent enhances its usefulness in adsorption applications.

5 As already noted, chemical modification can be used to alter the adsorptive properties of carbonaceous adsorbents. The range of chemical species which can be attached, however, is limited.

Bansal, Donnet and Stoeckli (in Chapter 5 of Active Carbon, Marcel Dekker, Inc., 1988) have reviewed different techniques of carbon surface modification. Physical
10 impregnation methods are described, as are methods that rely on chemical reactions with various species to modify the surface of the carbon. Some of the chemical surface modification techniques described by Bansal et al. are oxidation, halogenation, sulfonation, and ammoniation. Several of these techniques require treatment of the carbon at elevated temperatures. Another technique involving oxidation of the carbon with HNO_3
15 in the presence of a catalyst, has been described by Sircar and Golden (U.S. Patent No. 4,702,749). However, these techniques have certain disadvantages apparent to those familiar with the field.

In view of the limited utility of chemical means of modifying the surfaces of carbonaceous adsorbents, there still is a need to develop modified carbon adsorbents,
20 preferably with improved adsorption capacity over the unmodified material.

SUMMARY OF THE INVENTION

To achieve these and other advantages and in accordance with the purposes of the present invention, as embodied and broadly described herein, the present invention relates to an adsorbent composition containing a modified carbonaceous material capable of adsorbing an adsorbate.

Another embodiment of the present invention relates to a method to increase the adsorption capacity of a carbonaceous material capable of adsorbing an adsorbate or altering the adsorption isotherm of the adsorbate on the adsorbent, for instance, to allow an easier regeneration of the adsorbent. In this method, at least one organic group capable of increasing the adsorption capacity of a carbonaceous material is attached to the carbonaceous material.

The present invention, in addition, relates to a method of adsorbing an adsorbate and includes the step of contacting the adsorbate with a carbonaceous material which has been modified by attaching an organic group. The modified carbonaceous material is capable of adsorbing the adsorbate and at least one organic group is attached to the carbonaceous material.

Additional features and advantages of the present invention will be set forth in part in the description which follows, and in part will be apparent from the description, or may be learned by practice of the present invention. The objectives and other advantages of the present invention will be realized and attained by means of the elements and combinations particularly pointed out in the written description and appended claims.

It is to be understood that both the foregoing general description and the following detailed description are exemplary and explanatory only and are intended to provide further explanation of the present invention, as claimed.

The accompanying figures, which are incorporated in and constitute a part of this specification, illustrate several embodiments of the present invention and together with the description, serve to explain the principles of the present invention.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a graph plotting the amount of water adsorption on modified and unmodified carbon black.

Figure 2 is a graph plotting the amount of water adsorption on modified and unmodified activated carbon.

Figure 3 is a graph plotting the amount of water adsorption on modified and unmodified carbon black per unit surface area.

Figure 4 is a graph plotting the amount of water adsorption on modified and unmodified activated carbon per unit surface area.

Figure 5 is a graph plotting the amount of CO₂ adsorption on modified and unmodified carbon black at 273 K.

DETAILED DESCRIPTION OF THE PRESENT INVENTION

In further detail, an adsorbent composition of the present invention contains a modified carbonaceous material capable of adsorbing an adsorbate wherein at least one organic group is attached to the carbonaceous material.

5 The carbonaceous material capable of adsorbing an adsorbate includes, but is not limited to, activated carbon, carbon black, or other carbonaceous material obtained by the pyrolysis of cellulosic, fuel oil, polymeric, or other precursors. This includes carbonaceous material which is a waste product or by-product. Preferably, the carbonaceous material is activated carbon or carbon black capable of adsorbing an adsorbate. Commercial
10 examples of carbon black include, but are not limited to, Black Pearls® 2000 carbon black, Black Pearls® 430 carbon black, Black Pearls® 900 carbon black, and Black Pearls® 120 carbon black, all available from Cabot Corporation. Commercial examples of activated carbon include Darco S51, available from Norit; Sorbonorit 3, available from Norit; and BPL activated carbon from Calgon. ~~The~~ The carbonaceous material modified by
15 the procedures described herein may be a microporous or mesoporous activated carbon in granular or pellet form; a carbon black of different structures in fluffy or pelleted form; or any other carbonaceous material whose applicability to this invention is apparent to those skilled in the art, such as carbon fibers or carbon cloth. The choice of carbonaceous material used eventually depends on a variety of different factors, including
20 the application for which it is intended. Each of these types of carbonaceous material has the ability to adsorb at least one adsorbate. A variety of BET surface areas, micropore

volumes, and total pore volumes are available depending on the desired end use of the carbonaceous material.

Carbonaceous materials include, but are not limited to, material obtained by the compaction of small carbon particles and other finely divided forms of carbon as long as the carbon has the ability to adsorb at least one adsorbate and is capable of being chemically modified in accordance with the present invention.

~~The~~ The carbonaceous material described above is then modified by the attachment of an organic group to the carbonaceous material. Preferred processes for attaching an organic group to a carbonaceous material are described in detail in U.S. Patent Application Nos. 08/356,660, 08/572,525, 08/356,459, and 08/356,653, all incorporated in their entirety by reference herein. ~~These processes can be preferably used in preparing the modified carbon adsorbents of the present invention and permit the attachment of an organic group to the carbonaceous material via a chemical reaction.~~ As indicated above, the organic group attached to the carbonaceous material is one preferably capable of increasing the adsorption capacity of the carbonaceous material.

A preferred process for attaching an organic group to the carbonaceous materials involves the reaction of at least one diazonium salt with a carbonaceous material in the absence of an externally applied current sufficient to reduce the diazonium salt. That is, the reaction between the diazonium salt and the carbonaceous material proceeds without an external source of electrons sufficient to reduce the diazonium salt. Mixtures of different diazonium salts may be used. This process can be carried out under a variety

of reaction conditions and in any type of reaction medium, including both protic and aprotic solvent systems or slurries.

In another preferred process, at least one diazonium salt reacts with a carbonaceous material in a protic reaction medium. Mixtures of different diazonium salts may be used in this process. This process can also be carried out under a variety of reaction conditions.

Preferably, in both processes, the diazonium salt is formed *in situ*. If desired, in either process, the modified carbonaceous material can be isolated and dried by means known in the art. Furthermore, the modified carbonaceous material can be treated to remove impurities by known techniques. The various preferred embodiments of these processes are discussed below.

The processes can be carried out under a wide variety of conditions and in general are not limited by any particular condition. The reaction conditions must be such that the particular diazonium salt is sufficiently stable to allow it to react with the carbonaceous material. Thus, the processes can be carried out under reaction conditions where the diazonium salt is short lived. The reaction between the diazonium salt and the carbonaceous material occurs, for example, over a wide range of pH and temperature. The processes can be carried out at acidic, neutral, and basic pH. Preferably, the pH ranges from about 1 to 9. The reaction temperature may preferably range from 0°C to 100°C.

Diazonium salts, as known in the art, may be formed for example by the reaction of primary amines with aqueous solutions of nitrous acid. A general discussion of

diazonium salts and methods for their preparation is found in Morrison and Boyd, Organic Chemistry, 5th Ed., pp. 973-983, (Allyn and Bacon, Inc. 1987) and March, Advanced Organic Chemistry: Reactions, Mechanisms, and Structures, 4th Ed., (Wiley, 1992). According to this invention, a diazonium salt is an organic compound having one or more
5 diazonium groups.

The diazonium salt may be prepared prior to reaction with the carbonaceous material or, more preferably, generated *in situ* using techniques known in the art. *In situ* generation also allows the use of unstable diazonium salts such as alkyl diazonium salts and avoids unnecessary handling or manipulation of the diazonium salt. In particularly
10 preferred processes, both the nitrous acid and the diazonium salt are generated *in situ*.

A diazonium salt, as is known in the art, may be generated by reacting a primary amine, a nitrite and an acid. The nitrite may be any metal nitrite, preferably lithium
nitrite, sodium nitrite, potassium nitrite, or zinc nitrite, or any organic nitrite such as for
example isoamyl nitrite or ethyl nitrite. The acid may be any acid, inorganic or organic,
15 which is effective in the generation of the diazonium salt. Preferred acids include nitric acid, HNO₃, hydrochloric acid, HCl, and sulfuric acid, H₂SO₄.

The diazonium salt may also be generated by reacting the primary amine with an aqueous solution of nitrogen dioxide. The aqueous solution of nitrogen dioxide,
NO₂/H₂O, provides the nitrous acid needed to generate the diazonium salt.

20 Generating the diazonium salt in the presence of excess HCl may be less preferred than other alternatives because HCl is corrosive to stainless steel. Generation of the diazonium salt with NO₂/H₂O has the additional advantage of being less corrosive to

stainless steel or other metals commonly used for reaction vessels. Generation using $\text{H}_2\text{SO}_4/\text{NaNO}_2$ or $\text{HNO}_3/\text{NaNO}_2$ are also relatively non-corrosive.

In general, generating a diazonium salt from a primary amine, a nitrite, and an acid requires two equivalents of acid based on the amount of amine used. In an *in situ* process, the diazonium salt can be generated using one equivalent of the acid. When the primary amine contains a strong acid group, adding a separate acid may not be necessary. The acid group or groups of the primary amine can supply one or both of the needed equivalents of acid. When the primary amine contains a strong acid group, preferably either no additional acid or up to one equivalent of additional acid is added to a process of the invention to generate the diazonium salt *in situ*. A slight excess of additional acid may be used. One example of such a primary amine is para-aminobenzenesulfonic acid (sulfanilic acid).

In general, diazonium salts are thermally unstable. They are typically prepared in solution at low temperatures, such as $0-5^\circ\text{C}$, and used without isolation of the salt. Heating solutions of some diazonium salts may liberate nitrogen and form either the corresponding alcohols in acidic media or the organic free radicals in basic media.

However, the diazonium salt need only be sufficiently stable to allow reaction with the carbonaceous material. Thus, the processes can be carried out with some diazonium salts otherwise considered to be unstable and subject to decomposition. Some decomposition processes may compete with the reaction between the carbonaceous material and the diazonium salt and may reduce the total number of organic groups attached to the carbonaceous material. Further, the reaction may be carried out at

elevated temperatures where many diazonium salts may be susceptible to decomposition. Elevated temperatures may also advantageously increase the solubility of the diazonium salt in the reaction medium and improve its handling during the process. However, elevated temperatures may result in some loss of the diazonium salt due to other decomposition processes.

Reagents can be added to form the diazonium salt *in situ*, to a suspension of carbonaceous material in the reaction medium, *for example*, water. Thus, a carbonaceous material suspension to be used may already contain one or more reagents to generate the diazonium salt and the process accomplished by adding the remaining reagents.

Reactions to form a diazonium salt are compatible with a large variety of functional groups commonly found on organic compounds. Thus, only the availability of a diazonium salt for reaction with a carbonaceous material limits the processes of the invention.

The processes can be carried out in any reaction medium which allows the reaction between the diazonium salt and the carbonaceous material to proceed. Preferably, the reaction medium is a solvent-based system. The solvent may be a protic solvent, an aprotic solvent, or a mixture of solvents. Protic solvents are solvents, like water or methanol, containing a hydrogen attached to an oxygen or nitrogen and thus are sufficiently acidic to form hydrogen bonds. Aprotic solvents are solvents which do not contain an acidic hydrogen as defined above. Aprotic solvents include, for example, solvents such as hexanes, tetrahydrofuran (THF), acetonitrile, and benzonitrile. For a

discussion of protic and aprotic solvents see Morrison and Boyd, Organic Chemistry, 5th Ed., pp. 228-231, (Allyn and Bacon, Inc. 1987).

The processes are preferably carried out in a protic reaction medium, that is, in a protic solvent alone or a mixture of solvents which contains at least one protic solvent. Preferred protic media include, but are not limited to water, aqueous media containing water and other solvents, alcohols, and any media containing an alcohol, or mixtures of such media.

The reaction between a diazonium salt and a carbonaceous material can take place with any type of carbonaceous material, for example, in finely divided state or pelleted form. In one embodiment designed to reduce production costs, the reaction occurs during a process for forming carbonaceous material pellets. For example, a carbonaceous material product of the invention can be prepared in a dry drum by spraying a solution or slurry of a diazonium salt onto a carbonaceous material. Alternatively, the carbonaceous material product can be prepared by pelletizing a carbonaceous material in the presence of a solvent system, such as water, containing the diazonium salt or the reagents to generate the diazonium salt *in situ*. Aqueous solvent systems are preferred.

In general, the processes produce inorganic by-products, such as salts. In some end uses, such as those discussed below, these by-products may be undesirable. Several possible ways to produce a carbonaceous material product without unwanted inorganic by-products or salts are as follows:

First, the diazonium salt can be purified before use by removing the unwanted inorganic by-product using means known in the art. Second, the diazonium salt can be

generated with the use of an organic nitrite as the diazotization agent yielding the corresponding alcohol rather than an inorganic salt. Third, when the diazonium salt is generated from an amine having an acid group and aqueous NO_2 , no inorganic salts are formed. Other ways may be known to those of skill in the art.

5 In addition to the inorganic by-products, the process may also produce organic by-products. They can be removed, for example, by extraction with organic solvents. Other ways of obtaining products without unwarranted organic by-products may be known to those of skill in the art, and include washing or removal of ions by reverse osmosis.

10 The reaction between a diazonium salt and a carbonaceous material forms a carbonaceous material product having an organic group attached to the carbonaceous material. The diazonium salt may contain the organic group to be attached to the carbonaceous material. It may be possible to produce the carbonaceous material products of this invention by other means known to those skilled in the art.

15 The organic group may be an aliphatic group, a cyclic organic group, or an organic compound having an aliphatic portion and a cyclic portion. As discussed above, the diazonium salt employed can be derived from a primary amine having one of these groups and being capable of forming, even transiently, a diazonium salt. The organic group may be substituted or unsubstituted, branched or unbranched. Aliphatic groups include, for example, groups derived from alkanes, alkenes, alcohols, ethers, aldehydes, ketones, carboxylic acids, and carbohydrates. Cyclic organic groups include, but are not
20 limited to, alicyclic hydrocarbon groups (*for example*, cycloalkyls, cycloalkenyls), heterocyclic hydrocarbon groups (*for example*, pyrrolidinyl, pyrrolinyl, piperidinyl,

morpholinyl, and the like), aryl groups (*for example*, phenyl, naphthyl, anthracenyl, and the like), and heteroaryl groups (imidazolyl, pyrazolyl, pyridinyl, thienyl, thiazolyl, furyl, indolyl, and the like). As the steric hinderance of a substituted organic group increases, the number of organic groups attached to the carbonaceous material from the reaction between the diazonium salt and the carbonaceous material may be diminished.

When the organic group is substituted, it may contain any functional group compatible with the formation of a diazonium salt. Functional groups include, but are not limited to, R, OR, COR, COOR, OCOR, carboxylate salts such as COOLi, COONa, COOK, $\text{COO}^-\text{NR}_4^+$, halogen, CN, NR_2 , SO_3H , sulfonate salts such as SO_3Li , SO_3Na , SO_3K , $\text{SO}_3^-\text{NR}_4^+$, OSO_3H , OSO_3^- salts, $\text{NR}(\text{COR})$, CONR_2 , NO_2 , PO_3H_2 , phosphonate salts such as PO_3HNa and PO_3Na_2 , phosphate salts such as OPO_3HNa and OPO_3Na_2 , $\text{N}=\text{NR}$, NR_3^+X^- , PR_3^+X^- , S_xR , SSO_3H , SSO_3^- salts, $\text{SO}_2\text{NRR}'$, SO_2SR , SNRR' , SNQ , SO_2NQ , CO_2NQ , S-(1,4-piperazinediyl)-SR, 2-(1,3-dithianyl) 2-(1,3-dithiolanyl), SOR, and SO_2R . R and R', which can be the same or different, are independently hydrogen, branched or unbranched $\text{C}_1\text{-C}_{20}$ substituted or unsubstituted, saturated or unsaturated hydrocarbon, e.g., alkyl, alkenyl, alkynyl, substituted or unsubstituted aryl, substituted or unsubstituted heteroaryl, substituted or unsubstituted alkylaryl, or substituted or unsubstituted arylalkyl. The integer k ranges from 1-8 and preferably from 2-4. The anion X^- is a halide or an anion derived from a mineral or organic acid. Q is $(\text{CH}_2)_w$, $(\text{CH}_2)_x\text{O}(\text{CH}_2)_z$, $(\text{CH}_2)_x\text{NR}(\text{CH}_2)_z$, or $(\text{CH}_2)_x\text{S}(\text{CH}_2)_z$, where w is an integer from 2 to 6 and x and z are integers from 1 to 6. In the above formula, specific examples of R and R' are $\text{NH}_2\text{-C}_6\text{H}_4\text{-}$, $\text{CH}_2\text{CH}_2\text{-C}_6\text{H}_4\text{-NH}_2$, $\text{CH}_2\text{-C}_6\text{H}_4\text{-NH}_2$, and C_6H_5 .

Another example of an organic group is an aromatic group of the formula $A_y\text{Ar}$, which corresponds to a primary amine of the formula $A_y\text{ArNH}_2$. In this formula, the variables have the following meanings: Ar is an aromatic radical such as an aryl or heteroaryl group. Ar can be selected from the group consisting of phenyl, naphthyl, anthracenyl, phenanthrenyl, biphenyl, pyridinyl, benzothiadiazolyl, and benzothiazolyl; A is a substituent on the aromatic radical independently selected from a preferred functional group described above or A is a linear, branched or cyclic hydrocarbon radical (preferably containing 1 to 20 carbon atoms), unsubstituted or substituted with one or more of those functional groups; and y is an integer from 1 to the total number of -CH radicals in the aromatic radical. For instance, y is an integer from 1 to 5 when Ar is phenyl, 1 to 7 when Ar is naphthyl, 1 to 9 when Ar is anthracenyl, phenanthrenyl, or biphenyl, or 1 to 4 when Ar is pyridinyl.

Another set of organic groups which may be attached to the carbonaceous material are organic groups substituted with an ionic or an ionizable group as a functional group. An ionizable group is one which is capable of forming an ionic group in the medium of use. The ionic group may be an anionic group or a cationic group and the ionizable group may form an anion or a cation.

Ionizable functional groups forming anions include, for example, acidic groups or salts of acidic groups. The organic groups, therefore, include groups derived from organic acids. Preferably, when it contains an ionizable group forming an anion, such an organic group has a) an aromatic group or a C_1 - C_{12} alkyl group and b) at least one acidic group having a pKa of less than 11, or at least one salt of an acidic group having a pKa of less

than 11, or a mixture of at least one acidic group having a pKa of less than 11 and at least one salt of an acidic group having a pKa of less than 11. The pKa of the acidic group refers to the pKa of the organic group as a whole, not just the acidic substituent. More preferably, the pKa is less than 10 and most preferably less than 9. Preferably, the aromatic group or the C₁-C₁₂ alkyl group of the organic group is directly attached to the carbonaceous material. The aromatic group may be further substituted or unsubstituted, for example, with alkyl groups. The organic group can be a phenyl or a naphthyl group and the acidic group is a sulfonic acid group, a sulfinic acid group, a phosphonic acid group, or a carboxylic acid group. Examples of these acidic groups and their salts are discussed above. The organic group can be a substituted or unsubstituted sulfophenyl group or a salt thereof; a substituted or unsubstituted (polysulfo)phenyl group or a salt thereof; a substituted or unsubstituted sulfonaphthyl group or a salt thereof; or a substituted or unsubstituted (polysulfo)naphthyl group or a salt thereof. An example of a substituted sulfophenyl group is hydroxysulfophenyl group or a salt thereof.

Specific organic groups having an ionizable functional group forming an anion (and their corresponding primary amines for use in a process according to the invention) are p-sulfophenyl (p-sulfanilic acid), 4-hydroxy-3-sulfophenyl (2-hydroxy-5-amino-benzenesulfonic acid), and 2-sulfoethyl (2-aminoethanesulfonic acid).

Amines represent examples of ionizable functional groups that form cationic groups. For example, amines may be protonated to form ammonium groups in acidic media. Preferably, an organic group having an amine substituent has a pKb of less than 5. Quaternary ammonium groups (-NR₃⁺) and quaternary phosphonium groups

(-PR₃⁺) also represent examples of cationic groups. The organic group can contain an aromatic group such as a phenyl or a naphthyl group and a quaternary ammonium or a quaternary phosphonium group. The aromatic group is preferably directly attached to the carbonaceous material. Quaternized cyclic amines, and even quaternized aromatic amines, can also be used as the organic group. Thus, N-substituted pyridinium compounds, such as N-methyl-pyridyl, can be used in this regard. Examples of organic groups include, but are not limited to, (C₅H₄N)C₂H₅⁺X⁻, C₆H₄(NC₅H₅)⁺X⁻, C₆H₄COCH₂N(CH₃)₃⁺X⁻, C₆H₄COCH₂(NC₅H₅)⁺X⁻, (C₅H₄N)CH₃⁺X⁻, and C₆H₄CH₂N(CH₃)₃⁺X⁻, where X⁻ is a halide or an anion derived from a mineral or organic acid.

Aromatic sulfides encompass another group of organic groups. These aromatic sulfides can be represented by the formulas Ar(CH₂)_qS_k(CH₂)_rAr' or A-(CH₂)_qS_k(CH₂)_rAr'' wherein Ar and Ar' are independently substituted or unsubstituted arylene or heteroarylene groups, Ar'' is an aryl or heteroaryl group, k is 1 to 8 and q and r are 0-4. Substituted aryl groups would include substituted alkylaryl groups. Examples of arylene groups include phenylene groups, particularly p-phenylene groups, or benzothiazolylene groups. Aryl groups include phenyl, naphthyl and benzothiazolyl. The number of sulfurs present, defined by k preferably ranges from 2 to 4. Examples of carbonaceous material products are those having an attached aromatic sulfide organic group of the formula -(C₆H₄)_k-(C₆H₄)-, where k is an integer from 1 to 8, and more preferably where k ranges from 2 to 4. Other examples of aromatic sulfide groups are bis-para-(C₆H₄)₂-S₂-(C₆H₄)₂- and para-(C₆H₄)₂-S₂-(C₆H₅). The diazonium salts of these aromatic sulfide groups may be conveniently prepared from their corresponding primary amines, H₂N-Ar-S_k-Ar'-NH₂ or

$H_2N-Ar-S_k-Ar''$. Groups include dithiodi-4,1-phenylene, tetrathiodi-4,1-phenylene, phenyldithiophenylene, dithiodi-4,1-(3-chlorophenylene), $-(4-C_6H_4)-S-S-(2-C_6H_4NS)$, $-(4-C_6H_4)-S-S-(4-C_6H_4)-OH$, $-6-(2-C_6H_4NS)-SH$, $-(4-C_6H_4)-CH_2CH_2-S-S-CH_2CH_2-(4-C_6H_4)-$, $-(4-C_6H_4)-CH_2CH_2-S-S-S-CH_2CH_2-(4-C_6H_4)-$, $-(2-C_6H_4)-S-S-(2-C_6H_4)-$, $-(3-C_6H_4)-S-S-(3-C_6H_4)-$, $-6-(C_6H_4N_2S)$, $-6-(2-C_6H_4NS)-S-NRR'$ where RR' is $-CH_2CH_2OCH_2CH_2-$, $-(4-C_6H_4)-S-S-S-S-(4-C_6H_4)-$, $-(4-C_6H_4)-CH=CH_2$, $-(4-C_6H_4)-S-SO_3H$, $-(4-C_6H_4)-SO_2NH-(4-C_6H_4)-S-S-(4-C_6H_4)-NHSO_2-(4-C_6H_4)-$, $-6-(2-C_6H_4NS)-S-S-2-(6-C_6H_4NS)-$, $-(4-C_6H_4)-S-CH_2-(4-C_6H_4)-$, $-(4-C_6H_4)-SO_2-S-(4-C_6H_4)-$, $-(4-C_6H_4)-CH_2-S-CH_2-(4-C_6H_4)-$, $-(3-C_6H_4)-CH_2-S-CH_2-(3-C_6H_4)-$, $-(4-C_6H_4)-CH_2-S-S-CH_2-(4-C_6H_4)-$, $-(3-C_6H_4)-CH_2-S-S-CH_2-(3-C_6H_4)-$, $-(4-C_6H_4)-S-NRR'$, where RR' is $-CH_2CH_2OCH_2CH_2-$, $-(4-C_6H_4)-SO_2NH-CH_2CH_2-S-S-CH_2CH_2-NHSO_2-(4-C_6H_4)-$, $-(4-C_6H_4)-2-(1,3-dithianyl)$, and $-(4-C_6H_4)-S-(1,4-piperizinediyl)-S-(4-C_6H_4)-$.

Another set of organic groups which may be attached to the carbonaceous material are organic groups having an aminophenyl, such as $(C_6H_4)-NH_2$, $(C_6H_4)-CH_2-(C_6H_4)-NH_2$, $(C_6H_4)-SO_2-(C_6H_4)-NH_2$.

Any one or more of these organic groups, after attachment to the carbonaceous material which permits adsorption, and preferably an increase in the adsorption capacity of the carbonaceous material may be used in the present invention.

Preferably, the organic group attached to the carbonaceous material is an acid or base or a salt of an acid or base, and specific examples include phenyl or naphthyl groups having substituents like sulfonic acid, and carboxylic acid. Quaternary ammonium can also be used. Most preferred organic groups attached to the carbonaceous material are

(C₆H₄)-SO₃⁻Na⁺, (C₆H₄)-SO₃⁻K⁺, (C₆H₄)-SO₃⁻Li⁺, and the like. Generally, an acid-type organic group attachment will be useful in adsorbing basic adsorbates while a base-type organic group attachment will be useful in adsorbing acidic adsorbates.

5 A combination of different organic groups is possible. For instance, it is within the bounds of the present invention to attach more than one type of organic group to the same carbonaceous material or use a combination of carbonaceous materials, wherein some of the carbonaceous material has been modified with one organic group and another portion of the carbonaceous material has been modified with a different organic group. Varying degrees of modification are also possible, such as low weight percent or surface area modification, or a high weight percent or surface area modification. Also, mixtures of modified carbonaceous material and unmodified carbonaceous material can be used.

10 Preferably, the modified carbonaceous materials of the present invention, especially when the attached organic group is a phenyl or naphthyl group having substituents like sulfonic acid, carboxylic acid, or quaternary ammonium or salts thereof, can be directly analogous to polymeric ion exchange resins. These types of carbonaceous materials of the present invention can have one or more of the following properties as compared to conventional polymeric ion exchangers:

- 15 a) higher temperature stability;
- b) greater resistance to swelling; and
- 20 c) greater mechanical strength without adversely affecting uptake kinetics.

Furthermore, the modified carbonaceous materials of the present invention, besides being used as adsorbents, can also be used in separations ranging from water treatment

to metals separation/recovery, ion exchange, catalysis, and the like. An additional advantage of an adsorbent possessing exchangeable groups as described above is that it confers on the material the ability to be further surface modified using ion exchange procedures.

5 With respect to the adsorbates, any adsorbate capable of being adsorbed by one or more of the modified carbonaceous materials of the present invention is contemplated to be within the bounds of the present invention. Examples include, but are not limited to, polar species such as water, ammonia, mercaptans, sulfur dioxide, and hydrogen sulfide. By "polar species," it is understood that this is a species whose electronic structure is not symmetrical. This includes molecules that possess dipole moments, for example H₂O and NH₃; and/or molecules that possess quadrupole moments, such as CO₂ and molecules that possess unsaturated pi bonds (π), such as alkenes, alkynes, and other organic and inorganic compounds with double and triple bonds. Non-polar species such as argon, oxygen, methane, and the like can also be adsorbed with the appropriate modified carbonaceous materials of the present invention. In view of the description provided in this application, those skilled in the art will be able to determine which organic groups need to be attached to the carbonaceous materials in order to achieve the most effective adsorption affinity or increase in adsorption, depending upon the adsorbate and the adsorption processes involved.

20 By developing an adsorbent composition containing a modified carbonaceous material capable of adsorbing an adsorbate, selectivity for a particular adsorbate can be enhanced. Using the proper modified carbonaceous material, one can selectively adsorb

particular species from a multicomponent mixture. In other words, modifying the carbonaceous material to create the adsorbent composition of the present invention can decrease adsorption affinity for one component in order to maximize the adsorption affinity of another component which will maximize separation of the second component from the first component. Furthermore, by increasing adsorption of polar species, this further results in the relatively decreased adsorption of nonpolar species which improves selectivity. Further, the carbonaceous material can be modified in such a manner as to add a hydrophobic group to "disable" the oxygen functionalities on the surface of the carbonaceous material to increase the selectivity for the adsorption of nonpolar species.

The adsorbate can be in a liquid phase or in the gaseous or vapor phase, depending upon the needs and desires of the user. Certain adsorbates can be more efficiently adsorbed from the vapor or gaseous phases than from the liquid phase or vice versa, and the modified carbonaceous materials of the present invention are effective in adsorption from either phase.

One advantage of the present invention is to modify the surface of an activated carbon or carbon black adsorbent extensively, without damaging the structure or making the adsorbent more friable. For instance, a carbonaceous material can be surface modified based on the present invention with exchangeable sodium cations attached to the surface. This is very useful from the point of view of substituting different ions to alter the chemistry of the surface.

The beneficial effect of using the modified carbonaceous materials of the present invention for the purpose of adsorption can be demonstrated by comparing the adsorption

isotherms of an adsorbate on an unmodified carbonaceous adsorbent and the same carbonaceous adsorbent modified in accordance with the present invention.

The present invention will be further clarified by the following examples, which are intended to be purely exemplary of the present invention.

EXAMPLES

The effectiveness of the surface modification of exemplary carbonaceous material was determined by comparing the adsorption isotherms of various adsorbates on the unmodified carbonaceous materials, with adsorption isotherms on the carbonaceous materials modified in accordance with the present invention. Adsorbates used were water and CO₂, but other adsorbates could also be used.

Example 1

Pellets of Black Pearls® 430 carbon black and Darco S51 activated carbon from Norit were surface-modified using the following procedure:

Surface modification of Black Pearls® 430 Carbon Black:

A dispersion of 5 ml of water dilutable phenol-formaldehyde thermosetting resin (Schenectady International, Schenectady, NY) in 50 mL of water was mixed with 50 g of Black Pearls® 430 carbon black (available from Cabot Corp., Boston, MA). The mixture was pressed in 1 g portions using a 0.25 inch stainless steel die at a pressure of 5000 psi. The pellets were heated under flowing argon at 110°C for one hour and at 135°C for one hour to cure the resin. The temperature was then raised under flowing

argon at 20°C/min until a temperature of 650°C was reached. The temperature was then held at 650°C for three hours and cooled under flowing argon. The pellets were then crushed into pieces about 1 mm by 2 mm.

5 An aqueous solution of 0.81 g of sodium nitrite in about 1 g of water was added to a mixture of 16.8 g of the carbon black granules, 2.04 g of sulfanilic acid, and 50 g of water that was stirring at 84°C. After stirring for two hours, the resulting material was dried in an oven at 65°C.

Surface modification of activated carbon:

10 An aqueous solution of 30.5 g of sodium nitrite in about 100 g of water was added to a boiling mixture of 130 g of DARCO S51 activated carbon (available from Norit), 76.5 g of sulfanilic acid, and 1300 g of water. After stirring for 15 minutes, the heating was discontinued and the mixture was allowed to cool to room temperature with stirring. The resulting material was dried overnight in an oven at 50°C.

Ion Exchange:

15 The surface modified carbons were washed with a large amount of deionized water and dried. The material thus obtained was in the sodium form. Further modification of the carbon into potassium and lithium forms was carried out by ion exchange using 2M solutions of KOH and LiOH, respectively. The ion-exchanged material was washed thoroughly and dried. Adsorption experiments were carried out on the washed, dried
20 materials.

The surface areas of the unmodified and surface modified carbon materials are shown in Table 1 below. Surface areas were calculated from nitrogen (77 K) adsorption

data using the BET formalism (S.J. Gregg and K.S.W. Sing, in Adsorption, Surface Area, and Porosity, Academic Press, 1982). The adsorption experiments were carried out on an ASAP 2000 automated instrument, manufactured by Micromeritics Corp.

Table 1: Surface Areas and Pore Volumes of Unmodified and Surface Modified Materials

Sample ID	BET Surface Area, m ² /g	Pore Volume, cm ³ /g
BP 430 pellets, unmodified	99	0.518
BP 430, modified, washed	92	0.49
Darco S51	694	0.809
Darco S51, modified, washed	141.3	0.279

While the activated carbon lost some surface area and pore volume after the surface modification treatment, both the carbon black and the activated carbon underwent an increase in adsorption capacity per unit surface area as a result of the surface modification. The loss of any surface area and pore volume may be mitigated by pre-treating the carbonaceous material with immiscible organic solvent, like heptane. The results from adsorption of water vapor at 298 K on the unmodified and modified material are shown in Figure 1 (carbon black) and Figure 2 (activated carbon). The water adsorption experiments were carried out by a batch technique that involved equilibrating the sample with water vapor at a constant relative humidity, in a sealed cell. The constant relative humidities were attained by using saturated salt solutions, which have known relative humidities above their surface.

Both the activated carbon and carbon black contained Na^+ ions on the surface after the surface modification was carried out. The Na^+ ions can be substituted by other ions using standard ion exchange procedures (e.g., see Ion Exchange, by F. Helfferich, McGraw-Hill, 1962). The water adsorption isotherms for the surface-modified material with Na^+ ions on the surface, as well as the other ionic forms derived by ion exchange, are shown in Figures 1 and 2. The adsorption isotherms show the quantity of water vapor adsorbed, per gram of adsorbent, as a function of the relative pressure of water vapor. Figures 3 and 4 show the same data normalized by the BET surface area of the materials. It is clear that the adsorption capacities per unit area of the carbon black, Black Pearls® 430 carbon black and the Darco S51 carbon black, are considerably enhanced by the surface modification described in this invention. In addition, the shape of the water adsorption isotherm is changed as a result of the surface modification (concave upward, to linear or convex upward).

The surface modification technique of the present invention may affect the adsorption of gases like CO_2 as well, which possesses a quadrupole movement. Figure 5 shows the adsorption isotherm of CO_2 from the gas phase at 273 K on the same unmodified and modified Black Pearls® 430 carbon black. Adsorption of CO_2 was carried out on an ASAP 2000 automated adsorption system manufactured by Micromeritics Corp. The figure shows the quantity of CO_2 adsorbed as a function of the CO_2 pressure. Clearly, the adsorption of CO_2 is enhanced by the carbon surface modification technique described in this invention.

Other embodiments of the present invention will be apparent to those skilled in the art from consideration of the specification and practice of the invention disclosed herein. It is intended that the specification and examples be considered as exemplary only, with a true scope and spirit of the invention being indicated by the following claims.

WHAT IS CLAIMED IS:

1. An adsorbent composition comprising a modified carbonaceous material capable of adsorbing an adsorbate wherein at least one organic group is attached to said [modified] carbonaceous material,

5 2. The adsorbent composition of claim 1, wherein said organic group is $(C_6H_4)-SO_3^-Na^+$, $(C_6H_4)-SO_3^-Li^+$, or $(C_6H_4)SO_3^-K^+$.

3. The adsorbent composition of claim 1, wherein said organic group is $p-C_6H_4SO_2NH_2$ or $-C_6H_4NH_2$.

10 4. The adsorbent composition of claim 1, wherein said organic group is hydrophilic.

5. The adsorbent composition of claim 1, wherein said modified carbonaceous material is activated carbon or carbon black.

15 6. A method to increase the adsorption capacity of a carbonaceous material capable of adsorbing an adsorbate comprising attaching to said carbonaceous material at least one organic group capable of increasing said adsorption capacity.

7. The method of claim 6, wherein said carbonaceous material is activated carbon or carbon black.

8. The method of claim 6, wherein said organic group is $(C_6H_4)-SO_3^-Na^+$, $(C_6H_4)-SO_3^-Li^+$, or $(C_6H_4)SO_3^-K^+$.

5 9. The method of claim 6, wherein said organic group is $p-C_6H_4SO_2NH_2$ or $-C_6H_4NH_2$.

10. The method of claim 6, wherein said organic group is hydrophilic.

11. The method of claim 6, wherein said adsorbate is polar.

12. The method of claim 6, wherein said adsorbate is water, ammonia, carbon
10 dioxide, sulfur dioxide, or hydrogen sulfide.

13. The method of claim 12, wherein said adsorbate is water.

14. The method of claim 6, wherein said adsorbate is non-polar.

15. The method of claim 14, wherein said adsorbate is argon, oxygen, or methane.

16. The method of claim 6, wherein said adsorbate is in the gas or vapor phase.
17. The method of claim 6, wherein said adsorbate is in the liquid phase.
18. A method to adsorb an adsorbate comprising contacting said adsorbate with a modified carbonaceous material capable of adsorbing said adsorbate wherein at least one organic group is attached to said modified carbonaceous material.
19. The method of claim 18, wherein said modified carbonaceous material is activated carbon or carbon black.
20. The method of claim 18, wherein said organic group is $(\text{C}_6\text{H}_4)\text{-SO}_3\text{Na}^+$, $(\text{C}_6\text{H}_4)\text{-SO}_3\text{Li}^+$, or $(\text{C}_6\text{H}_4)\text{SO}_3\text{K}^+$.
21. The method of claim 18, wherein said organic group is $\text{p-C}_6\text{H}_4\text{SO}_2\text{NH}_2$ or $\text{-C}_6\text{H}_4\text{NH}_2$.
22. The method of claim 18, wherein said organic group is hydrophilic.
23. The method of claim 18, wherein said adsorbate is polar.

24. The method of claim 18, wherein said adsorbate is water, ammonia, carbon dioxide, hydrogen sulfide, argon, oxygen, or methane.

25. The method of claim 24, wherein said adsorbate is water.

26. The adsorbent composition of claim 1, wherein said organic group contains an exchangeable ion.

27. An ion exchange material comprising the adsorbent composition of claim 26.

ABSTRACT OF THE DISCLOSURE

An adsorbent composition containing a modified carbonaceous material capable of adsorbing an adsorbate is disclosed, wherein at least one organic group is attached to the carbonaceous material. Furthermore, methods to increase the adsorption capacity of a carbonaceous material capable of adsorbing an adsorbate and methods to adsorb an adsorbate using the above-described adsorbent composition are also disclosed.

5

Figure 1: Water vapor adsorption on unmodified Black Pearls 430 carbon black and on various ionic forms of modified material

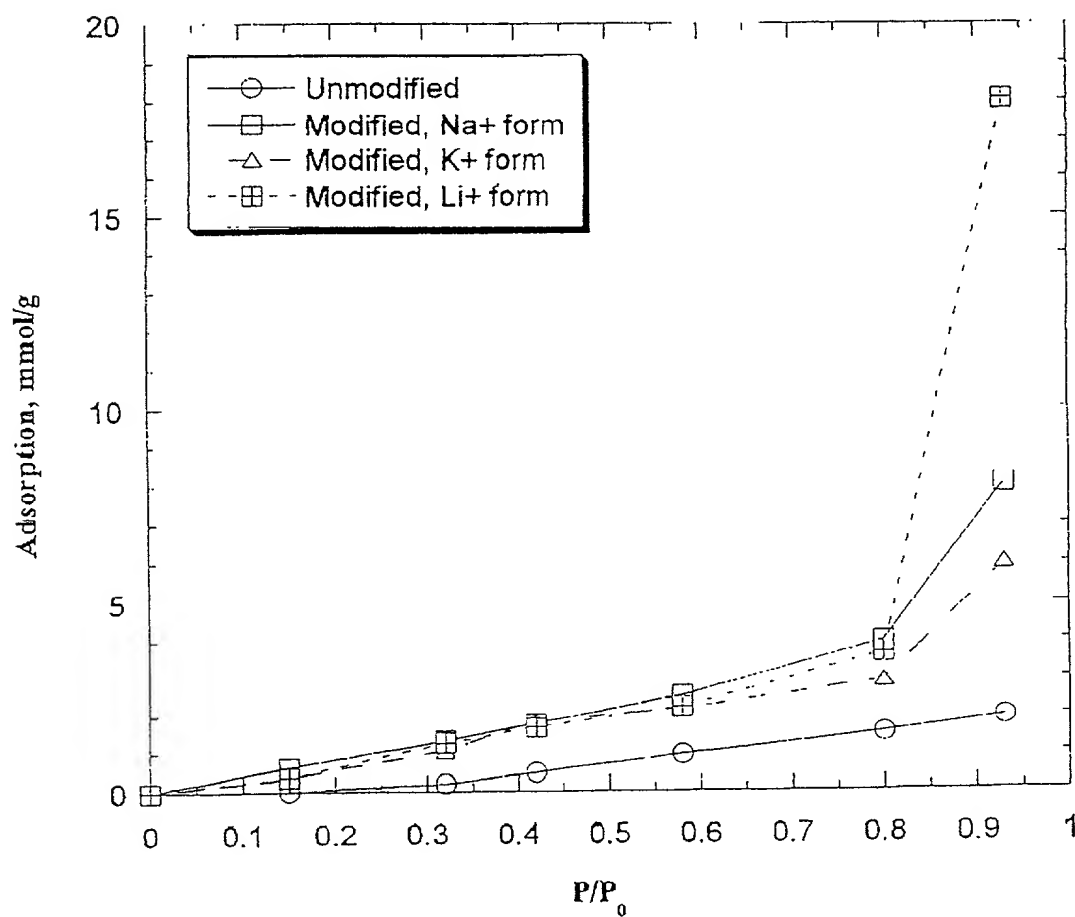


Figure 2: Water vapor adsorption on unmodified Darco S51 activated carbon and on modified ionic forms

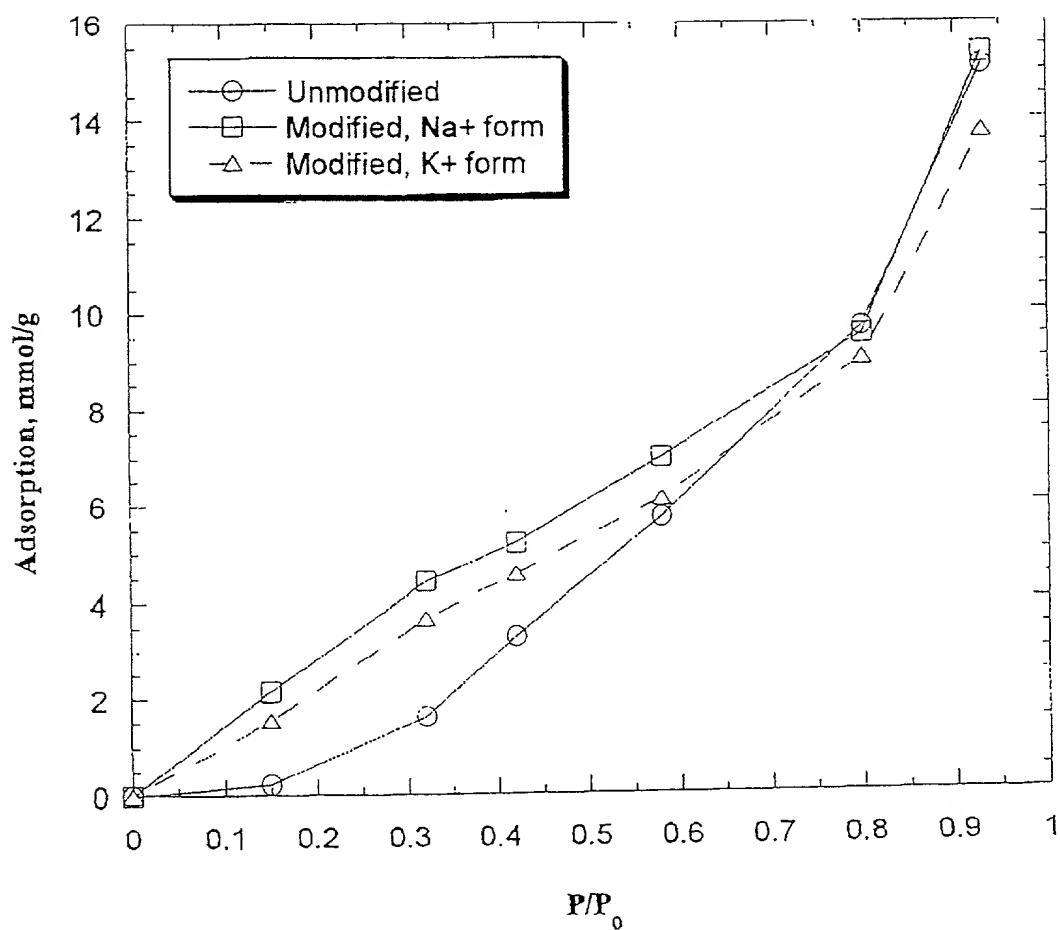


Figure 3: Water vapor adsorption per unit surface area for unmodified and modified Black Pearls 430 carbon black

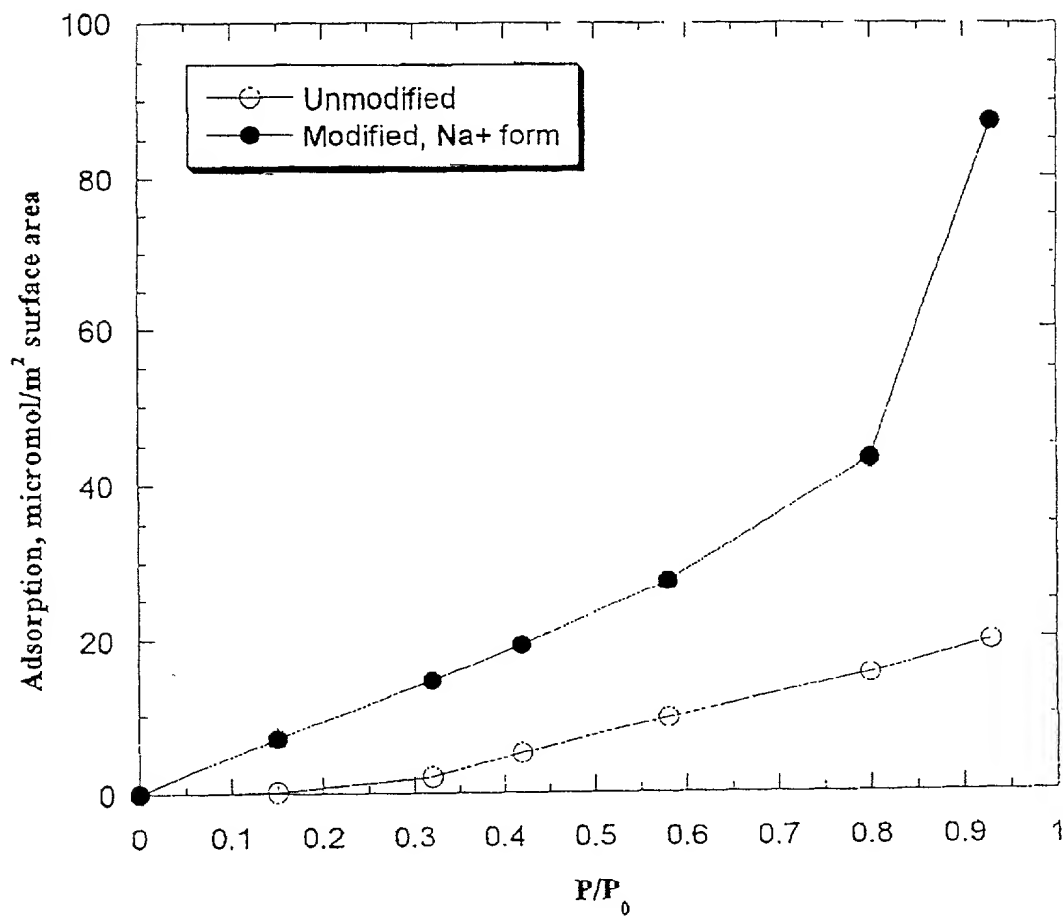


Figure 4: Water vapor adsorption per unit surface area for unmodified and modified Darco S51 activated carbon

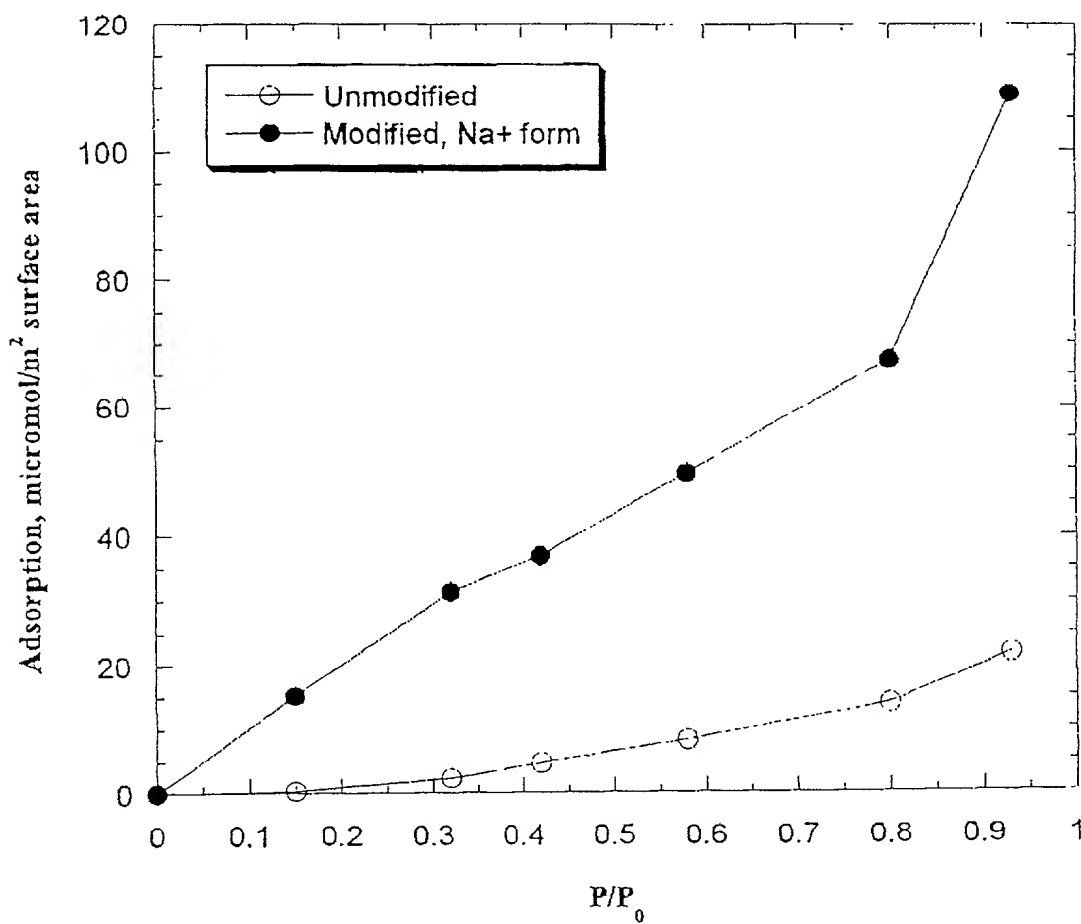
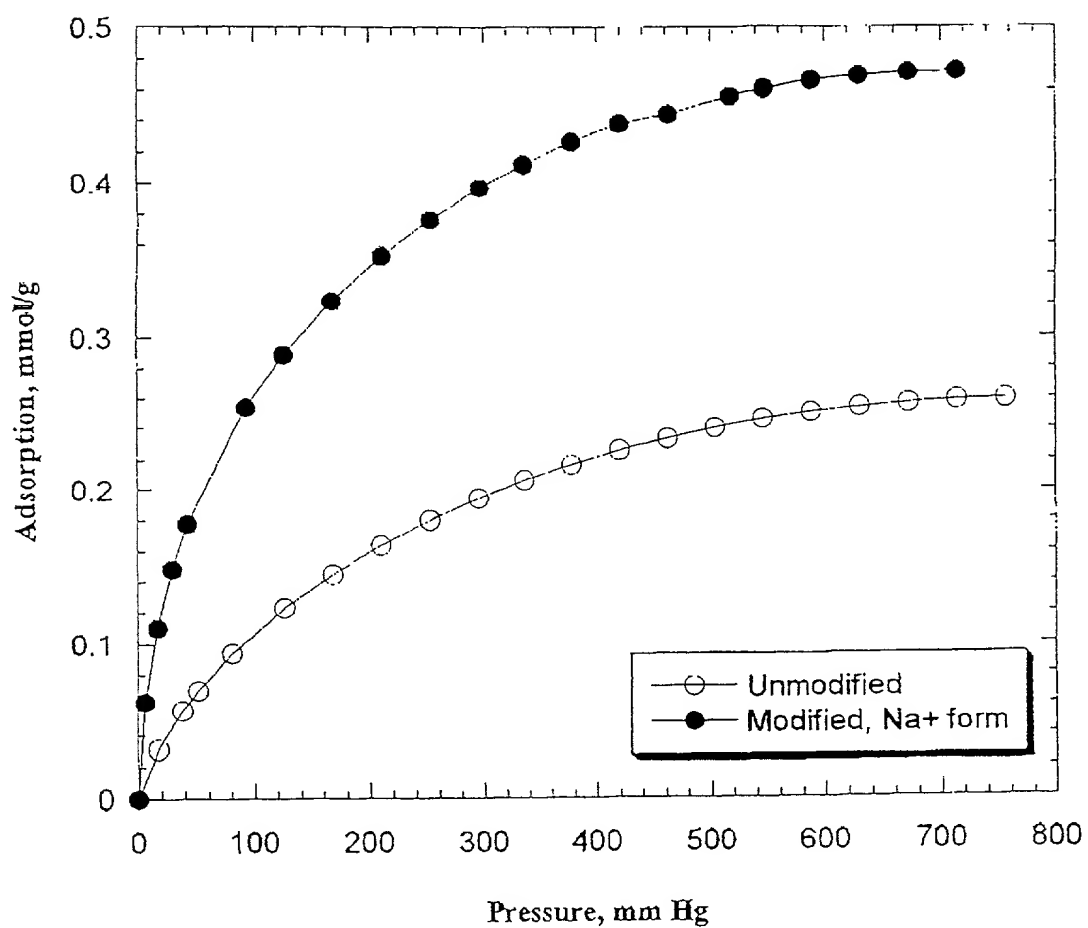


Figure 5: Adsorption of CO_2 at 273 K on unmodified and modified Black Pearls 430 carbon black



DECLARATION AND POWER OF ATTORNEY

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below my name;

I believe I am an original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled " **Modified Carbon Adsorbents And Processes For Adsorption Using The Same,**" the specification of which [] is attached and/or [x] was filed as Serial No. 08/663,709 on June 14, 1996 and was amended on _____ (if applicable); or was filed as PCT International Application Number _____ on _____ and was amended on _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, §1.56(a).

I hereby claim foreign priority benefits under Title 35, United States Code, §119 of any foreign applications for patent or inventor's certificate or of any PCT international application(s) designating at least one country other than the United States of America listed below and have also identified below any foreign application(s) for patent or inventor's certificate or any PCT international application(s) designating at least one country other than the United States of America filed by me on the same subject matter having a filing date before that of the application(s) of which priority is claimed:

COUNTRY (If PCT indicate PCT)	APPLICATION NO.	DATE OF FILING (day, month, year)	PRIORITY CLAIMED UNDER 35 USC §119
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) or PCT international application(s) designating the United States of America that is/are listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in that/those prior application(s) in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, § 1.56(a) which occurred between the filing

Declaration and Power of Attorney

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date of the prior application(s) and the national or PCT international filing date of this application:

U.S. APPLICATIONS		STATUS (Check one)		
U.S. APPLICATION NO.	U.S. FILING DATE	PATENTED	PENDING	ABANDONED
PCT APPLICATIONS DESIGNATING THE U.S.				
PCT APPLICATION NO.	PCT FILING DATE	U.S. SERIAL NO. ASSIGNED (if any)		

I hereby appoint Martha Ann Finnegan (Registration No. 31,453), Michelle B. Lando (Registration No. 33,941), Clair X. Mullen, Jr. (Registration No. 20,348) and Luke A. Kilyk (Registration No. 33,251) as my attorney with full power of substitution and revocation, to prosecute this application and transact all business in the United States Patent and Trademark Office connected therewith.

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Billerica, MA 01821

and direct all telephone calls to Martha Ann Finnegan at (508) 670-6198.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the

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Citizen

Attorney Docket No. 96074

PATENT

United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Inventor's Signature: _____

Full Name of First Inventor: Ranjan Ghosal

Date: ____ August, 1996

Residence: 1429 Columbia Drive, NE, Albuquerque, New Mexico 87106

Post Office Address: 1429 Columbia Drive, NE, Albuquerque, New Mexico 87106

Citizenship: India

Inventor's Signature: James A. Belmont

Full Name of Second Inventor: James A. Belmont

Date: 14 August, 1996

Residence: 8 Conant Street, Acton, Massachusetts 01720

Post Office Address: 8 Conant Street, Acton, Massachusetts 01720

Citizenship: United States of America

Inventor's Signature: _____

Full Name of First Inventor: Douglas M. Smith

Date: ____ August, 1996

Residence: 1412 Marquette P1, NE, Albuquerque, New Mexico, 87106

Post Office Address: 1412 Marquette P1, NE, Albuquerque, New Mexico, 87106

Citizenship: United States of America

Declaration and Power of Attorney

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Attorney Docket No. 96074

PATENT

Attorney

Inventor's Signature: Jameel Menashi

Full Name of First Inventor: Jameel Menashi

Date: 14 August, 1996

Residence: 68 Gleason Road, Lexington, Massachusetts 02173

Post Office Address: 68 Gleason Road, Lexington, Massachusetts 02173

Citizenship: United States of America

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DECLARATION AND POWER OF ATTORNEYdate of
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As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below my name;

I believe I am an original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled " **Modified Carbon Adsorbents And Processes For Adsorption Using The Same,**" the specification of which [] is attached and/or [] was filed as Serial No. 08/663,709 on June 14, 1996 and was amended on _____ (if applicable); or was filed as PCT International Application Number _____ on _____ and was amended on _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, §1.56(a).

I hereby claim foreign priority benefits under Title 35, United States Code, §119 of any foreign applications for patent or inventor's certificate or of any PCT international application(s) designating at least one country other than the United States of America listed below and have also identified below any foreign application(s) for patent or inventor's certificate or any PCT international application(s) designating at least one country other than the United States of America filed by me on the same subject matter having a filing date before that of the application(s) of which priority is claimed:

COUNTRY (If PCT indicate PCT)	APPLICATION NO.	DATE OF FILING (day, month, year)	PRIORITY CLAIMED UNDER 35 USC §119
			[] Yes [] No
			[] Yes [] No
			[] Yes [] No
			[] Yes [] No
			[] Yes [] No
			[] Yes [] No

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) or PCT international application(s) designating the United States of America that is/are listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in that/those prior application(s) in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, § 1.56(a) which occurred between the filing

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date of the prior application(s) and the national or PCT international filing date of this application:

U.S. APPLICATIONS		STATUS (Check one)		
U.S. APPLICATION NO.	U.S. FILING DATE	PATENTED	PENDING	ABANDONED
PCT APPLICATIONS DESIGNATING THE U.S.				
PCT APPLICATION NO.	PCT FILING DATE	U.S. SERIAL NO. ASSIGNED (if any)		

I hereby appoint Martha Ann Finnegan (Registration No. 31,453), Michelle B. Lando (Registration No. 33,941), Clair X. Mullen, Jr. (Registration No. 20,348) and Luke A. Kilyk (Registration No. 33,251) as my attorney with full power of substitution and revocation, to prosecute this application and transact all business in the United States Patent and Trademark Office connected therewith.

Please address all communications regarding this application to:

Martha Ann Finnegan, Esq.
Cabot Corporation
157 Concord Road
Billerica, MA 01821

and direct all telephone calls to Martha Ann Finnegan at (508) 670-6198.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the

Attorney:
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Inventor:

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Post Office

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Attorney Docket No. 96074

PATENT

United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Inventor's Signature: _____

Full Name of First Inventor: Ranjan Ghosal

Date: ____ August, 1996

Residence: 1429 Columbia Drive, NE, Albuquerque, New Mexico 87106

Post Office Address: 1429 Columbia Drive, NE, Albuquerque, New Mexico 87106

Citizenship: India

Inventor's Signature: _____

Full Name of Second Inventor: James A. Belmont

Date: ____ August, 1996

Residence: 8 Conant Street, Acton, Massachusetts 01720

Post Office Address: 8 Conant Street, Acton, Massachusetts 01720

Citizenship: United States of America

Inventor's Signature: _____

Full Name of First Inventor: Douglas M. Smith

Date: 17 August, 1996

Residence: 1412 Marquette P1, NE, Albuquerque, New Mexico, 87106

Post Office Address: 1412 Marquette P1, NE, Albuquerque, New Mexico, 87106

Citizenship: United States of America

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DECLARATION AND POWER OF ATTORNEY

As a below named inventor, I hereby declare that:

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COUNTRY (if PCT indicate PCT)	APPLICATION NO.	DATE OF FILING (day, month, year)	PRIORITY CLAIMED UNDER 35 USC §119
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) or PCT international application(s) designating the United States of America that is/are listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in that/those prior application(s) in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, § 1.56(a) which occurred between the filing

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date of the prior application(s) and the national or PCT international filing date of this application:

U.S. APPLICATIONS		STATUS (Check one)		
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I hereby appoint Martha Ann Finnegan (Registration No. 31,453), Michelle B. Lando (Registration No. 33,941), Clair X. Mullen, Jr. (Registration No. 20,348) and Luke A. Kilyk (Registration No. 33,251) as my attorney with full power of substitution and revocation, to prosecute this application and transact all business in the United States Patent and Trademark Office connected therewith.

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Attorney Docket No. 96074

PATENT

United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Inventor's Signature: Ranjan Ghosal

Full Name of First Inventor: Ranjan Ghosal

Date: 15 August, 1996

Residence: 1429 Columbia Drive, NE, Albuquerque, New Mexico 87106

Post Office Address: 1429 Columbia Drive, NE, Albuquerque, New Mexico 87106

Citizenship: India

Inventor's Signature: _____

Full Name of Second Inventor: James A. Belmont

Date: ____ August, 1996

Residence: 8 Conant Street, Acton, Massachusetts 01720

Post Office Address: 8 Conant Street, Acton, Massachusetts 01720

Citizenship: United States of America

Inventor's Signature: _____

Full Name of First Inventor: Douglas M. Smith

Date: ____ August, 1996

Residence: 1412 Marquette Pl, NE, Albuquerque, New Mexico, 87106

Post Office Address: 1412 Marquette Pl, NE, Albuquerque, New Mexico, 87106

Citizenship: United States of America

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- 3 -

Attorney

Inventor

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PATENT

Full Name of First Inventor: Jameel Menashi

Residence: 68 Gleason Road, Lexington, Massachusetts 02173

Post Office Address: 68 Gleason Road, Lexington, Massachusetts 02173

Citizenship: United States of America

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Declaration and Power of Attorney